

Baseline tissue Doppler imaging-derived echocardiographic parameters and left ventricle reverse remodelling following cardiac resynchronization therapy introduction

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Abstract

Introduction: The aim of the study was to assess the relation of baseline mechanical dyssynchrony with the left ventricular end-systolic volume (LVESV) decrease following cardiac resynchronization (CRT) therapy introduction.

Material and methods: Sixty consecutive patients (aged 66.3 ± 8.7 years; 57 men) with chronic heart failure (71.7% of ischaemic and 28.3% of non-ischaemic origin) and current indications for CRT were assessed before and 3 months after biventricular heart stimulator implantation. Longitudinal movements of twelve segments of the left ventricle (LV) (6 basal and 6 midlevel) and two segments of the right ventricle (RV) were analysed using tissue Doppler imaging (TDI) techniques with time from onset of Q wave in ECG to peak systolic velocity in colour-coded TDI (T_{TDI}), time to peak strain (T_{strain}) and time to peak strain rate ($T_{strain\ rate}$). Minimal and maximal time differences within LV and between LV and RV walls were calculated.

Results: In the study group LVEF and 6-min walk test distance increased, while NYHA class, NT-proBNP level, left ventricular end-diastolic volume and LVESV decreased. Significant correlations between the magnitude of LVESV reduction with maximal time differences between T_{strain} of 12 LV segments ($r = 0.34$, $p = 0.017$) and time differences between T_{TDI} basal LV-RV segments ($r = -0.29$, $p = 0.041$) were found.

Conclusions: Only a few TDI-derived parameters such as maximal time differences between T_{strain} of 12 LV segments and T_{TDI} difference of LV-RV basal segments can be useful to predict the magnitude of left ventricle reverse remodelling after CRT introduction.

Key words: heart failure, pacemakers, echocardiography, Doppler dyssynchrony.

Introduction

Cardiac resynchronization therapy (CRT) has a well-established position in the treatment of drug-refractory chronic heart failure (CHF) [1, 2]. The CRT improves filling of the left ventricle, increases ejection fraction (LVEF), cardiac output and systolic blood pressure, reduces mitral insufficiency and, in long-term observations, induces left ventricle reverse remodelling. The CRT was also found to decrease sleep-related breathing disorders that

augment CHF, improve the function of the autonomic nervous system and lead to a fall of some inflammatory markers [3-6]. Solid evidence from clinical trials has proven that CRT amends physical capacity and quality of life, and decreases morbidity and mortality related to CHF. Unfortunately, a considerable proportion (30-40%) of individuals undergoing CRT in clinical trials did not benefit from this novel therapy [7]. Since the absence of echocardiographic dyssynchrony is associated with a less favourable CRT outcome, the answer to the question of CRT failure is also believed to be found in the complex heart mechanical performance [8]. A debate on the issue has been triggered with echocardiographic researchers from around the world contributing volubly to the discussion.

The aim of this study was to assess the relation of baseline mechanical dyssynchrony in the echocardiographic study with the use of tissue Doppler imaging (TDI), strain and strain rate parameters with the left ventricle reverse remodelling observed after CRT implantation, a powerful long-term prognostic factor, in typical daily practice CHF patients receiving biventricular heart stimulation [9, 10]. A unique approach in the analysis of CRT outcome was engaged: the presence of at least 10% relative left ventricular end-systolic volume (LVESV) reduction or the magnitude of LVESV decrease following biventricular heart stimulator was considered.

Material and methods

Study group

Seventy-two consecutive patients with chronic heart failure in NYHA class III-IV stable for at least 3 months despite optimized pharmacotherapy, with left ventricle end-diastolic diameter (LVEDd) > 55 mm, left ventricular ejection fraction (LVEF) ≤ 35% and wide QRS complex (≥ 120 ms), were involved in the prospective study. Eight individuals met the exclusion criteria, two refused to have CRT introduced, and in another two implantation of the left ventricular lead did not succeed. Eventually 60 patients (aged 66.3 ± 8.7 years, 57 men [95%], 3 women [5%]) participated in the study.

Exclusion criteria

Exclusion criteria comprised the presence of unstable angina, acute myocardial infarction, coronary artery bypass graft or percutaneous coronary intervention within 3 months; continuous or intermittent intravenous inotropic drug therapy, an estimated life expectancy of less than 12 months, a mechanical right-side heart valve, heart transplant, pregnancy or concurrent enrolment in a study thought to confound the results.

Study design

Before CRT implantation and at 3 months of the follow-up (12-16 weeks) individuals were evaluated clinically with NYHA class, 6-min walk test (6-MWT), history with questions about hospitalizations and echocardiographically. Criteria of diagnosis of different clinical and biochemical disorders were adopted from the European Society of Cardiology (ESC) guidelines [1]. All patients undergoing CRT had had coronarography done. Ischaemic background of CHF was diagnosed when there was at least 50% stenosis of one or more coronary artery branches or a patient had a history of coronary artery bypass graft (CABG) or percutaneous coronary intervention (PCI) [11, 12].

Informed consent was obtained from each patient. The study protocol conforms to the ethical guidelines of the 1975 Declaration of Helsinki and was accepted by the Jagiellonian University Medical College Bioethical Committee.

Biventricular device implantation

Patients in sinus rhythm received an atrio-biventricular stimulator (DDDR BiV – $n = 47$), while individuals with permanent atrial fibrillation received a biventricular device (VVIR BiV – $n = 13$). Patients with a history of cardiac arrest and/or malignant ventricular arrhythmias had the system with combined cardioverter-defibrillator (CRT-D) implanted (18 cases). During CRT all leads were implanted transvenously. The left ventricular lead, guided by a venogram, was placed in a coronary sinus tributary in a stable lateral or postero-lateral position, with a < 3.5 V capture threshold. The right ventricular lead was placed in the septal or outflow tract (RVOT) position. Positions of lead tips were verified on frontal and sagittal chest X-ray. Two patients required left ventricular lead repositioning due to their dislocation. Atrioventricular (AV) delay remained standard programmed 150/120 ms, unless conduction of cardiotoxic systoles was preserved. In those patients AV was shortened until ventricles were paced (5 cases). Interventricular (VV) timing left nominal 5 ms, unless no signs of biventricular stimulation in the body surface ECG were observed. VV was changed then to elicit the picture of QRS fusion beats in ECG lead V1 (7 cases) like in the study of Bailey *et al.* [13]. Such a VV-delay optimization was there proven to correlate better with maximum rate of rise of left ventricular pressure (dp/dt_{max}) than the strategy to obtain the narrowest paced QRS complexes.

In patients with permanent atrial fibrillation, ventricles' rate control with β -blockers and digoxin was assessed. In one patient an unsatisfactory pharmacologic effect (< 95% of biventricular stimulation) was observed and ablation of the

atrioventricular junction was performed. The follow-up was prolonged over 3 months after the procedure.

Echocardiography

Echocardiographic examinations were performed with the apparatus Vivid 7 (General Electric Healthcare). Left ventricular volumes and LVEF were evaluated using biplane Simpson's method. All stored echo recordings were analysed off-line by means of the Echo Pack system with GE brand software by an experienced physician with respect to the echocardiography norms of the European Association of Echocardiography and recommendations of the American Society of Echocardiography (ASE) for performance and reporting of echocardiographic studies for CRT [14, 15]. A different approach from the ASE recommendations regarded four aspects. Patients with atrial fibrillation were included – analysis of atrial fibrillation was done on at least 5-beat sequences of 60-80 bpm rate. The search for the desired peak TDI, strain rate and strain was not limited to the parts of curves within the time when the aortic valve was open. Multippeak curves with peaks of similar height were excluded from the analysis. Each single myocardial wall was recorded separately (not in sets of two opposite walls) in order to record longitudinal segments' movements along scan lines and reduce the Doppler angle dependent bias.

Longitudinal movements of twelve segments of the left ventricle (LV) (6 basal and 6 midlevel) and two segments of the right ventricle (RV) were analysed using tissue Doppler imaging (TDI) techniques with times from onset of Q wave in ECG to peak systolic velocity in colour-coded TDI (T_{TDI}), time to peak strain (T_{strain}) and time to peak strain rate ($T_{strain\ rate}$). Standard deviations (SD) of times of LV segments were calculated, and minimum and maximum time differences among LV segments as well as LV and RV walls in these different techniques were analysed.

The following dyssynchrony indices were calculated:

- 1) T-SD – standard deviation of time from the beginning of QRS complex to maximum systolic velocity/strain/strain rate among 12 left ventricle segments,
- 2) T-12 – maximum difference of time from the beginning of QRS complex to maximum systolic velocity/strain/strain rate among 12 left ventricle segments (basal and midlevel),
- 3) T-6-basal – maximum difference of time from the beginning of QRS complex to maximum systolic velocity/strain/strain rate among 6 basal left ventricle segments,
- 4) T-IVS- free wall – maximum difference of time from the beginning of QRS complex to maximum systolic velocity/strain/strain rate between basal and midlevel septal segments (septal and antero-septal segments) and left ventricle free wall (lateral and posterior segments),
- 5) T-basal IVS-free wall – maximum difference of time from the beginning of QRS complex to maximum systolic velocity/strain/strain rate between basal septal segments (septal and antero-septal segments) and left ventricle free wall (lateral and posterior segments),
- 6) T-IVD – maximum difference of time from the beginning of QRS complex to maximum systolic velocity/strain/strain rate between basal left ventricle and right ventricle segments.

Echocardiographic recordings interpretation limitations

Poor image quality and the ambiguity of interpretation of the obtained curves caused that 26.4% of TDI, strain rate and strain parameters were incalculable. Based on the analysis of 10 randomly chosen recordings, intra-observer variability for LVESV estimation was $4 \pm 3\%$, for peak systolic velocity – $7 \pm 10\%$, for peak systolic strain – $9 \pm 12\%$ and for peak systolic strain rate – $9 \pm 11\%$.

The CRT outcome assessment

Patients with at least 10% relative reduction of LVESV after 3 months of the follow-up were classified as responders to CRT. Clinical and echocardiographic parameters were compared between these two groups. In the second part of the analysis correlations between TDI-derived variables and the magnitude of reverse remodelling as a CRT outcome measure were appraised.

Statistical analysis

Statistical analysis was performed within SAS System 9.1 (SAS Institute Inc., Cary North Carolina, USA) by Student's *t*-test, Fisher's exact test and χ^2 test. The PROC CORR procedure was applied to assess correlations. Statistical significance was considered when $p < 0.05$. Data are expressed as mean (\pm SD) for continuous variables and as absolute frequencies or relative percentages for categorical variables.

Results

Clinical characteristics of the study group

In the study group 61.6% were in sinus rhythm, 16.7% had paroxysmal atrial fibrillation and 21.7% permanent fibrillation. Conduction disorders constituted left bundle branch block (65.0%), right bundle branch block (1.7%), and ventricular

conduction disturbances of other morphology (33.3%). Two patients with previously implanted ventricular heart stimulator (VVI), 9 with dual chamber pacemaker (DDD) and 3 with dual chamber implantable cardioverter-defibrillator (DDD-ICD) had their systems upgraded to CRT. Clinical characteristics of the study group are summarized in Table I.

Course of the study

Three patients died in the 3-month-follow-up: one of stroke, one of myocardial infarction, and one suddenly at home of unknown cause. These individuals were excluded from the further analysis. In the observation period 10 patients were hospitalized, 7 of them due to CHF exacerbation. All prescriptions (medications – see Table I) remained stable through the study.

Effects of CRT

In the whole study group LVEF increased, 6-MWT distance rose, and left ventricular end-diastolic volume (LVEDV) and LVESV decreased. Mean NYHA class and QRS width diminished (Table II). Results of specialized examinations before and after

implementation in the studied subgroups of responders ($n = 34$, 59.7%) and non-responders to CRT ($n = 23$, 40.3%) are shown in Table III.

Clinical and classic echocardiographic features

Responders did not differ from non-responders in respect of baseline left ventricular end-diastolic and end-systolic diameters and volumes, or the size of mitral regurgitation (Table III).

Baseline dyssynchrony echocardiographic indices and CRT outcome

In the analysis of colour-coded TDI, myocardial strain rate and strain indices, the only parameter that differentiated the groups with and without reverse remodelling was T_{strain} of 12 LV segments (Table IV). Significant correlations between the magnitude of LVESV reduction with maximal time differences between T_{strain} of 12 LV segments and time differences between T_{TDI} basal LV-RV segments were found (Table V, Figure 1).

Multivariate analysis

In the multivariate analysis, variables that differentiated the groups with and without $\geq 10\%$

Table I. Baseline clinical data of the whole study group and of responders and non-responders subgroups according to the applied criterion of at least 10% end-systolic left ventricular volume reduction at 3-months of follow-up (data presented as mean value with standard deviation (SD) or number of patients with the percentage share (%) of the whole analysed group or subgroup, p – value for comparison between responders and non-responders)

Parameter	All ($n = 60$)	Responders ($n = 34$)	Non-responders ($n = 23$)	Value of p
Age [years]	66.3 (8.7)	65.8 (8.2)	67.1 (9.4)	0.60
Gender – male [%]	57 (95%)	33 (97%)	24 (92%)	0.57
BMI [kg/m ²]	26.1 (4.3)	26.0 (3.0)	27.0 (5.0)	0.33
NYHA	3.1 (0.2)	3.1 (0.3)	3.1 (0.2)	0.91
Ischaemic aetiology of CHF	43 (71%)	21 (62%)	22 (84%)	0.04
Permanent atrial fibrillation	13 (22%)	6 (18%)	7 (27%)	0.048
Diabetes type 2	25 (42%)	12 (35%)	13 (50%)	0.30
Chronic obstructive pulmonary disease	11 (18%)	7 (21%)	4 (15%)	0.52
Hypercholesterolaemia	46 (76%)	26 (76%)	20 (76%)	0.99
Anaemia	2 (3%)	2 (6%)	0 (0%)	0.35
CRT-D	18 (30%)	11 (32%)	7 (27%)	0.56
History of stroke	4 (6.7%)	3 (9%)	1 (3%)	0.62
Chronic renal disease	20 (33.3%)	8 (23%)	12 (46%)	0.09
Medication				
ACEI/ARB	51 (85%)	30 (88%)	21 (81%)	0.48
β -Blocker	58 (96.7%)	33 (97%)	25 (96%)	0.99
Loop diuretic	53 (88.3%)	28 (82%)	25 (96%)	0.12
Potassium-sparing diuretic	40 (66.7%)	25 (73%)	15 (58%)	0.27
Digoxin	13 (21.7%)	8 (23%)	5 (19%)	0.76
Amiodarone	18 (30%)	9 (26%)	9 (35%)	0.39

CRT-D – cardiac resynchronization therapy defibrillator, ACEI – angiotensin converting enzyme inhibitor, ARB – angiotensin receptor blocker

Table II. Clinical and echocardiographic parameters in the study group before and 3 months after CRT introduction

Parameter	Before CRT (n = 57)	3 months after CRT (n = 57)	Value of p
NYHA	3.1 (0.2)	2.2 (0.7)	< 0.001
6-MWT [m]	298.0 (107.4)	373.1 (127.1)	< 0.001
LVEDd [mm]	73.3 (8.9)	71.5 (9.8)	0.003
LVESd [mm]	62.4 (10.0)	60.6 (11.3)	0.034
LVEDV [ml]	244.3 (83.8)	226.4 (88.6)	0.002
LVESV [ml]	192.8 (71.9)	168.7 (76.5)	< 0.001
SV [ml]	51.5 (16.6)	57.7 (16.9)	< 0.001
LVEF [%]	21.7 (4.8)	26.0 (4.8)	< 0.001
NT-proBNP [pg/ml]	2387.2 (1836.1)	1844.5 (1312.3)	0.027
QRS width	184.2 (28.3)	152.7 (19.1)	< 0.001

NYHA – New York Heart Association functional class, 6-MWT – 6-min walk test distance, LVEDd – left ventricular end-diastolic diameter, LVESd – left ventricular end-systolic diameter, LVEDV – left ventricular end-diastolic volume, LVESV – left ventricular end-systolic volume, SV – stroke volume, LVEF – left ventricular ejection fraction, NT-proBNP – N-terminal pro-B-type natriuretic peptide

Table III. Specialized examinations results of responders and non-responders according to the applied criterion of at least 10% end-systolic left ventricular volume reduction at 3-month follow-up

Parameter	Responders, n = 34 (59.6%)			Non-responders, n = 23 (40.4%)		
	Initially	After 3 months	Value of p	Initially	After 3 months	Value of p
LVEDd [mm]	73.2 (7.4)	69.9 (8.3)	< 0.001	73.5 (10.9)	73.9 (11.5)	0.47
LVESd [mm]	62.0 (7.9)	58.1 (9.4)	0.003	63.0 (12.7)	64.2 (13.0)*	0.13
LVEDV [ml]	246.3 (61.1)	209.5 (58.7)	< 0.001	241.3 (110.7)	251.4 (117.6)	0.054
LVESV [ml]	194.5 (55.9)	153.0 (51.4)	< 0.001	190.3 (92)	191.8 (99.9)	0.68
SV [ml]	51.6 (12.8)	56.3 (12.4)	0.002	51.4 (21.3)	59.6 (22.2)	0.002
MR [%]	25.6 (14)	18.4 (11)	0.001	23.7 (11)	24.4 (11)	0.62
QRS [ms]	179 (28)	153 (20)	0.001	199 (27)	152 (17)	0.001
LVEF [%]	21.5 (4.9)	27.1 (4.8)	< 0.001	21.9 (4.7)	24.5 (4.7)*	0.003
6-MWT [m]	315.7 (112.1)	392.3 (109.5)	< 0.001	271.9 (96)	344.7 (147.4)	0.001
NYHA	3.1 (0.3)	2.1 (0.5)	< 0.001	3.1 (0.2)	2.5 (0.8)	0.001
NT-proBNP [pg/ml]	2005.6 (1632.3)	1512.0 (1148.7)	0.012	3150.5 (2034.3)#	2509.5 (1403.6)*	0.32
VV-sep [cm]		8.1 (2.9)			8.9 (2.3)	

6-MWT – 6-min walk test distance, LVEDd – left ventricular end-diastolic diameter, LVEDV – left ventricular end-diastolic volume, LVEF – left ventricular ejection fraction, LVESd – left ventricular end-systolic diameter, LVESV – left ventricular end-systolic volume, MR – mitral regurgitation – presented as % of left atrium area, NT-proBNP – N-terminal pro-B-type natriuretic peptide, NYHA – New York Heart Association Functional Classification, VV-sep. – right and left ventricular electrode tips' separation on sagittal chest X-ray
Each parameter is presented as arithmetic mean with standard deviation (SD), p – for the difference between initial and after 3-month follow-up values of the same parameter, #p < 0.05 – for the difference between initial parameters of responders and non-responders, *p < 0.05 – for the difference between parameters of responders and non-responders measured after 3 months of follow-up

LVESV reduction in the follow-up were included: an ischaemic aetiology of CHF, permanent atrial fibrillation, prevalence of hypertension, hypotension and N-terminal pro-B-type natriuretic peptide (NT-proBNP) concentration [16]. Also left bundle branch block (LBBB) and non-LBBB morphology of QRS complex were considered. The predictive values of T_{strain} of 12 LV segments and T_{TDI} basal LV-RV regarding LVESV reduction lost their statistical significance.

Discussion

Significance of response to CRT criteria choice. Rationale for the magnitude of LVESV reduction assessment as CRT outcome measure

Response to CRT criteria hugely influence study results. Moreover, response to CRT rates differ dramatically depending on the applied definition. Compiling data from the literature suggest that reversal of the maladaptive left ventricle remodelling

Table IV. Baseline parameters [ms] of mechanical dyssynchrony assessed with colour-coded tissue Doppler imaging (TDI), myocardial strain and strain rate in the whole study group, responders and non-responders according to the applied criterion of at least 10% end-systolic left ventricular volume reduction at 3-month follow-up

Parameter	All	Responders	Non-responders	Value of p
Colour-coded TDI				
T _{TDI} -SD	58.0 (16)	54.8 (18)	62 (12.8)	0.14
T _{TDI} -12	176.2 (54)	165.8 (62)	189.1 (41)	0.15
T _{TDI} -6-basal	153.2 (96)	159.5 (111)	143.8 (66)	0.55
T _{TDI} -IVS-free wall	222.8 (181)	200 (180)	258.3 (183)	0.29
T _{TDI} -basal IVS-free wall	130.9 (88)	126.8 (85)	137.4 (96)	0.69
T _{TDI} -IVD	59.4 (46)	49.5 (40)	73.7 (51)	0.06
Strain				
T _{strain} -SD	125.2 (33)	131.83 (30)	115.6 (36)	0.09
T _{strain} -12	414.6 (111)	441.9 (114)	367.7 (89)	0.023
T _{strain} -6-basal	318.9 (109)	342.2 (108)	287.5 (106)	0.09
T _{strain} -IVS-free wall	391.2 (243)	405.9 (228)	495.3 (269)	0.60
T _{strain} -basal IVS-free wall	210.7 (139)	229.0 (141)	180 (135)	0.22
T _{strain} -IVD	169.3 (103)	161.5 (100)	179.5 (108)	0.56
Strain rate				
T _{strain rate} -SD	133.4 (29)	140.6 (27)	124.7 (31)	0.06
T _{strain rate} -12	403.8 (98)	418.3 (100)	384.1 (94)	0.22
T _{strain rate} -6-basal	341.9 (78)	359.6 (68)	318 (87)	0.07
T _{strain rate} -IVS-free wall	454.7 (283)	494.1 (306)	395.5 (242)	0.25
T _{strain rate} -basal IVS-free wall	243.6 (144)	271.7 (141)	192.5 (140)	0.08
T _{strain rate} -IVD	169.2 (116)	177.5 (124)	158.6 (108)	0.58

T-SD – standard deviation of time from the beginning of QRS complex to maximum systolic velocity/strain/strain rate in 12 left ventricle segments, T-12 – maximum difference of time from the beginning of QRS complex to maximum systolic velocity/strain/strain rate among 12 left ventricle segments (basal and midlevel), T-6-basal – maximum difference of time from the beginning of QRS complex to maximum systolic velocity/strain/strain rate among 6 basal left ventricle segments, T-IVS-free wall – maximum difference of time from the beginning of QRS complex to maximum systolic velocity/strain/strain rate between basal and midlevel septal segments (septal and antero-septal segments) and left ventricle free wall (lateral and posterior segments), T-basal IVS-free wall – maximum difference of time from the beginning of QRS complex to maximum systolic velocity/strain/strain rate between basal septal segments (septal and antero-septal segments) and left ventricle free wall (lateral and posterior segments), T-IVD – maximum difference of time from the beginning of QRS complex to maximum systolic velocity/strain/strain rate between basal left ventricle and right ventricle segments

mechanism in CHF observed a few months after CRT implementation is the best predictor of positive long-term outcome. Yu *et al.* showed in a series of 141 CRT patients that reduction of LVESV of at least 10% was related to a very low event rate (6.9% all-cause mortality) in a mean follow-up period of 695 days [9]. Di Biase *et al.* in an analysis from a prospective registry including 398 consecutive patients, with the median duration of follow-up of 4.4 years, demonstrated that the magnitude of reverse remodelling measured by LVEF and LVESV after 3 months of CRT implementation irrespectively of aetiology predicted outcome [10].

Methodological considerations

Our analyses of the colour-coded TDI velocities, strain and strain rate curves were not restrained to

the ejection period. This common practice distorts dyssynchrony imaging since contraction of certain left ventricle parts takes place not only during the ejection phase but also before opening and after the aortic valve closure. Such velocities may be greater than the ejection peaks and need to be distinguished from passive tethering by the adjoining myocardial segments. Both these movements are interchangeably classified by different authors as the phenomenon of post-systolic shortening/thickening (PSS). The PSS is therefore commonly observed in patients undergoing CRT, was found to be of high sensitivity in detecting ischaemia in other studies, but might be present even in one third of left ventricle segments in healthy individuals [18]. A problem also arises when fine tuning of the myocardial region of interest during TDI analysis fails to produce a single

reproducible peak. Strategies of proceeding in such situations varied in different studies, and the ASE recommendation of choosing the earlier peak if two or more peaks are of the same height is also challenged. In our research multi-peak curves with comparable maximal height were excluded from the analysis. As described in the 'material and methods' section, using the quality experience of previous studies, the offline calculations were performed by one experienced physician with the variability of the studied TDI, strain and strain rate parameters not exceeding 10%, which has to be considered as an attribute of this paper [19, 20].

One of our study assumptions was to provide information useful for typical CHF patients undergoing CRT. It is crucial to remember that, although presence of permanent atrial fibrillation is believed to predict poorer outcome, the huge meta-analysis of Upadhyay *et al.* of 1,164 CRT individuals indicates that the strategy, undertaken also in our study, of ventricles' rate control with β -blockers, digoxin and amiodarone and, if ineffective, ablation of the atrioventricular node confers a benefit similar to that of CRT to the group of patients in sinus rhythm [2, 21]. Despite reservations about the accuracy of echocardiographic analysis in atrial fibrillation patients, some researchers appreciate the significance of mechanical dyssynchrony in this group and successfully apply its quantification in their studies [22, 23].

Efficiency of colour-coded TDI derived indices of baseline dyssynchrony and interventricular dyssynchrony appraisal

The largest body of literature and the principal method in clinical use to quantify mechanical dyssynchrony in CHF patients undergoing CRT is

Table V. Correlation between the magnitude of LVESV reduction [ml] following CRT introduction and initial parameters of mechanical dyssynchrony [ms] assessed with colour-coded tissue Doppler imaging (TDI), myocardial strain and strain rate in the whole study group

Parameter	Value of <i>r</i>	Value of <i>p</i>
Colour-coded TDI		
T _{TDI} -SD	-0.05	0.72
T _{TDI} -12	-0.07	0.63
T _{TDI} -6-basal	-0.03	0.80
T _{TDI} -IVS-free wall	-0.16	0.28
T _{TDI} -basal IVS-free wall	-0.04	0.76
T _{TDI} -IVD	-0.29	0.041
Strain		
T _{strain} -SD	0.18	0.24
T _{strain} -12	0.34	0.017
T _{strain} -6-basal	0.21	0.16
T _{strain} -IVS-free wall	0.15	0.30
T _{strain} -basal IVS-free wall	0.07	0.64
T _{strain} -IVD	0.05	0.76
Strain rate		
T _{strain rate} -SD	0.19	0.19
T _{strain rate} -12	0.20	0.16
T _{strain rate} -6-basal	0.14	0.34
T _{strain rate} -IVS-free wall	0.15	0.33
T _{strain rate} -basal IVS-free wall	0.08	0.60
T _{strain rate} -IVD	0.03	0.81

Abbreviations – see Table IV

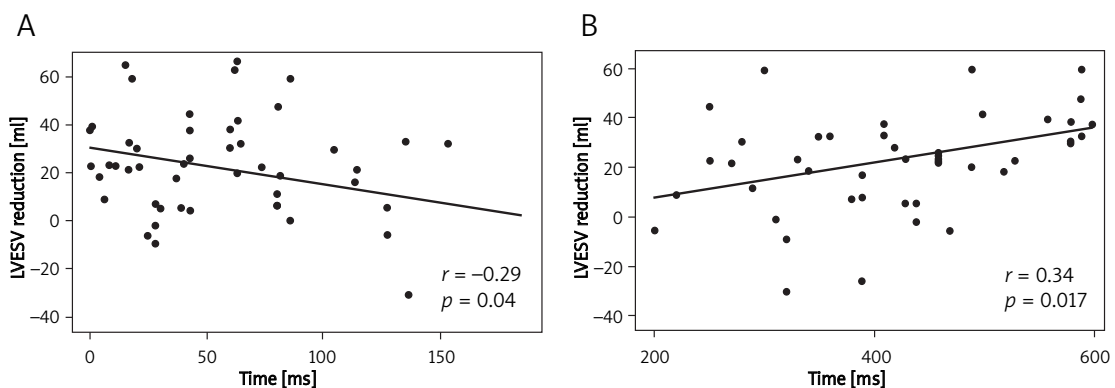


Figure 1. Correlation between the magnitude of LVESV reduction following CRT introduction and initial parameters of mechanical dyssynchrony: **A** – T_{TDI} basal LV-RV segments, **B** – T_{strain} of 12 LV segments

T_{TDI} basal LV-RV segments – maximum difference of time from the beginning of QRS complex to maximum systolic velocity between basal left ventricle and right ventricle segments, T_{strain}-12 – maximum difference of time from the beginning of QRS complex to maximum systolic strain among 12 left ventricle segments

represented by the assessment of longitudinal left ventricle shortening velocities using colour-coded TDI from the apical window [15]. We found no significant correlation of the analysed colour-coded TDI parameters of intraventricular dyssynchrony with LVESV reduction. Similar results come from other studies on correlations of TDI baseline indices with LVESV volume changes after CRT. Miyazaki *et al.* failed to demonstrate the usefulness of septal-lateral, anteroseptal-posterior delay and T_{TDI-SD} in a median 6-month-follow-up [24]. Phillips *et al.* found no predictive value of septal-lateral delay [25].

The interventricular dyssynchrony assessed with maximal difference between time from the beginning of QRS complexes to the peak velocity of basal LV and RV segments correlated negatively, indicating that the smaller the initial difference the more pronounced the reverse remodelling after CRT, which is quite an interesting finding. The echocardiographic assessment of interventricular dyssynchrony in previous studies on CRT was mainly based on the interventricular mechanical delay index (IVMD). The IVMD, reproducible and easily calculated in daily practice had little efficacy in predicting response to CRT as compared to intraventricular dyssynchrony assessment [26-29]. It should also be noted that evaluation of interventricular dyssynchrony with TDI and evaluation with IVMD represent different phenomena of heart performance: TDI, the movement of heart muscle parts; IVMD, the blood flow. The timing of the ejection of blood out of ventricles is conditioned by many factors including valves' and right ventricle function, different haemodynamic states, pulmonary diseases, etc., and poorly reflects the synchrony of myocardial contraction. The role of interventricular dyssynchrony and the optimal approach to the issues of TDI parameters are yet to be clarified.

Advantage of myocardial strain calculation. Strain rate imaging inapplicable

Strain and strain rate imaging, thought to be unaffected by movement of structures in the thorax, adjoining parts of heart muscle or the location in the ventricle, have the theoretic advantage of differentiating active myocardial contraction from passive translational movement [30]. Unfortunately, they are difficult to calculate in spherical LV geometry common in CHF, their application requires proper training and experience of the operator as well as good quality images, and they have disputable reproducibility [15].

Advantages of strain imaging in the prediction of CRT outcome were emphasized by some researchers [31-33]. Breithardt *et al.* concluded that the degree of dyssynchrony was not accurately represented by the timing of myocardial velocity,

especially in ischaemic heart disease, and proposed that the timing of deformation should be the preferred modality [31]. Sogaard *et al.* found that the extent of delayed longitudinal contraction at the base predicted LVEF increase following CRT [32, 33]. In our study the only strain parameter that differentiated responders from non-responders and correlated with the magnitude of LVESV reduction was $T_{strain-12}$. In the aforementioned study of Miyazaki, baseline $T_{strain-SD}$ showed a significant correlation with the percent reduction in LVESV at the follow-up ($r = 0.57, p < 0.001$) [24]. This supports the idea of eliciting the sites of the earliest and the latest activation with the strain imaging technique.

Parameters of strain rate in our study were not related to the change of LVESV. This is coherent with the ASE report based on the series of the studies of Yu *et al.*, who failed to demonstrate the usefulness of strain rate imaging to predict the LV reverse remodelling response to CRT [34, 35].

The multivariate analysis showed that the predictive significance of dyssynchrony appraisal is lost when other variables are taken into account. This emphasizes the role of complex evaluation of potential CRT recipients including a broad array of conditions associated with poor prognosis in CHF [1].

Our study group, after inclusion of patients with atrial fibrillation and the group of pacemakers upgraded to CRT, was heterogeneous but highly reflected the clinical profile of patients undergoing biventricular stimulator implantation in daily practice. Recording myocardial walls separately reduced Doppler angle dependent bias in calculating echocardiographic parameters, but caused that analysed curves, though always at least 3 were considered, came from different heart cycles.

The study had a non-randomized design, the sample size was small and the observational period was relatively short. Clinical and heart performance changes elicited by CRT may not always appear simultaneously and patients who present with clinical effects may not exhibit echocardiographic improvement and vice versa; thus longer follow-up would definitely provide accessory information about the significance of left ventricle reverse remodelling noted 3 months after CRT introduction [36].

In conclusion, only a few TDI-derived parameters such as maximum time difference from the beginning of QRS complex to maximum strain among 12 basal and midlevel left ventricular segments or maximum time difference from the beginning of QRS complex to maximum myocardium velocity in colour-coded TDI between left and right ventricular basal segments can be useful to predict the magnitude of left ventricle reverse remodelling after CRT introduction. The predictive value of the parameters is limited when other clinical and biochemical factors are considered.

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